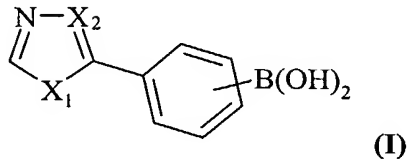


**CLAIMS**

1. A process for the preparation of a compound of the Formula I



5 wherein,

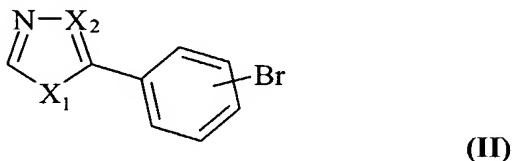
$X_1$  is selected from O,  $NR_1$  or S; and

$X_2$  is selected from CH or N;

wherein  $R_1$  is a nitrogen-protecting group,

which comprises :-

- 10 the sequential reaction of a compound of the Formula II



with,

- (i) methyl- or an optionally substituted aryl- lithium; and then  
 (ii) n-butyl-, s-butyl-, t-butyl- or n-hexyl- lithium; and then  
 15 (iii) a borate ester.

2. The process according to claim 1 wherein  $X_1$  is O.

3. The process according to claim 1 or 2 wherein  $X_2$  is N.

20

4. The process according to any one of claims 1-3 wherein said methyl- or an optionally substituted aryl- lithium is 4-methylphenyllithium or methyllithium.

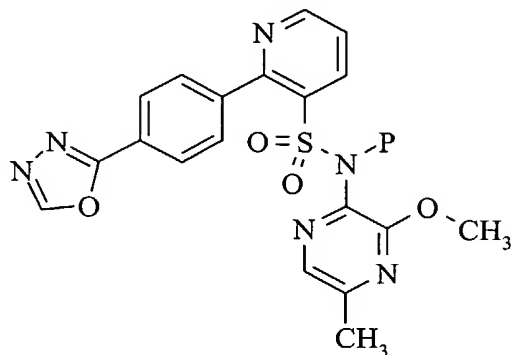
5. The process according to any one of claims 1-4 wherein said n-butyl-, s-butyl-,  
 25 t-butyl- or n-hexyl- lithium is n-hexyllithium or n-butyllithium.

6. The process according to any one of claims 1-5 wherein said borate ester is triisopropylborate.

- 16 -

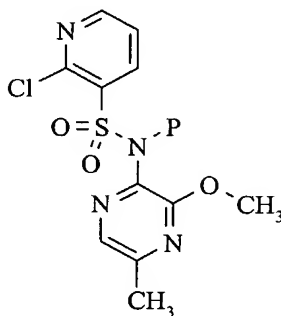
7. [4-(1,3,4-Oxadiazol-2-yl)phenyl]boronic acid prepared by the process as claimed in any one of claims 1-6.

8. A process for preparing compounds of Formula IV:



(IV)

which comprises coupling [4-(1,3,4-oxadiazol-2-yl)phenyl]boronic acid with a compound of Formula III:



(III)

wherein P is a nitrogen-protecting group.

10

9. The process according to claim 8 which takes place in the presence of

(i) a source of palladium (0) selected from PdCl<sub>2</sub>, Pd(Ph<sub>3</sub>P)<sub>4</sub> or Pd(OAc)<sub>2</sub>;

(ii) a suitable ligand selected from triphenylphosphine or 3,3',3''-phosphinidyne tris(benzenesulphonic acid) trisodium salt;

15 (iii) a base selected from triethylamine, benzyldimethylamine, N-methylmorpholine, N-methylpiperidine, triethanolamine, ethyldiethanolamine, diisopropylethylamine, potassium acetate, cesium fluoride or potassium fluoride.

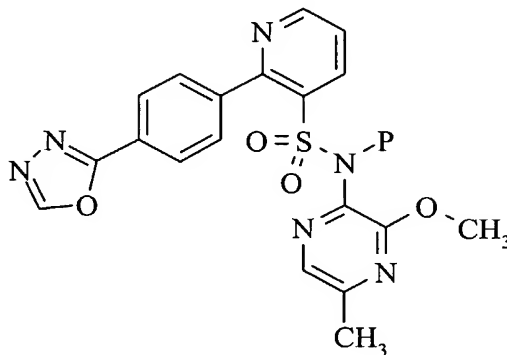
- 17 -

10. The process according to claim 8 or claim 9 wherein said [4-(1,3,4-oxadiazol-2-yl)phenyl]boronic acid is prepared according to the process as claimed in any one of claims 1-7.

11. The process according to any one of claims 8 - 10 wherein P is isobutoxycarbonyl.

5

12. A compound of Formula IV:



(IV)

wherein P is a nitrogen-protecting group.

10 13. A compound of Formula IV as claimed in claim 11 which is N-(isobutoxycarbonyl) N-(3-methoxy-5-methylpyrazin-2-yl)-2-(4-[1,3,4-oxadiazol-2-yl]phenyl) pyridine-3-sulphonamide.